

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶: A23L 2/44, 2/39	A1	(11) International Publication Number: WO 96/22704 (43) International Publication Date: 1 August 1996 (01.08.96)
(21) International Application Number: PCT/JP96/00109 (22) International Filing Date: 22 January 1996 (22.01.96) (30) Priority Data: 7/28804 24 January 1995 (24.01.95) JP (71) Applicant (for all designated States except US): OTSUKA PHARMACEUTICAL CO., LTD. [JP/JP]; 9, Kandatsukasa-cho 2-chome, Chiyoda-ku, Tokyo 101 (JP). (72) Inventors; and (75) Inventors/Applicants (for US only): TAKAICHI, Akihisa [JP/JP]; 172-3, Aza Nakajima, Takashima, Narutocho, Naruto-shi, Tokushima 772 (JP). OKAMOTO, Toshihiko [JP/JP]; 632, Ko, Kokufucho, Tokushima-shi, Tokushima 772 (JP). FUKUDA, Tetsuo [JP/JP]; Mezon-Kitahama 401, 21-9, Aza Miyano Higashi, Kitahama, Muiyacho, Naruto-shi, Tokushima 772 (JP). (74) Agent: KAMEI, Hirokatsu; Ai Association of Patent and Trade-mark Attorneys, 12F, Kinyukoko-Sumitomoseimei Building, 5-20, Minamihommachi 4-chome, Chuo-ku, Osaka-shi, Osaka 541 (JP).		(81) Designated States: AU, CA, CN, KR, US, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE). Published <i>With international search report.</i>
(54) Title: STABLE PRESERVATION METHOD OF POWDERED SOFT DRINK PREPARATION AND POWDERED SOFT DRINK PREPARATION (57) Abstract The present invention provides a stable preservation method of a powdered soft drink preparation, which comprises adding 0.2 to 1.0 % by weight of a calcium oxide and 0.02 to 2.0 % by weight of a particulate silicon dioxide in the powdered soft drink preparation containing a carbohydrate and a sour agent as a main component. The powdered soft drink preparation of the present invention maintains the original powder state, color and taste, which are obtained immediately after the production, for a long period of time without causing comparatively rapid coagulation of powders, discoloration and change in taste at the time of preservation.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AM	Armenia	GB	United Kingdom	MW	Malawi
AT	Austria	GE	Georgia	MX	Mexico
AU	Australia	GN	Guinea	NE	Niger
BB	Barbados	GR	Greece	NL	Netherlands
BE	Belgium	HU	Hungary	NO	Norway
BF	Burkina Faso	IE	Ireland	NZ	New Zealand
BG	Bulgaria	IT	Italy	PL	Poland
BJ	Benin	JP	Japan	PT	Portugal
BR	Brazil	KE	Kenya	RO	Romania
BY	Belarus	KG	Kyrgyzstan	RU	Russian Federation
CA	Canada	KP	Democratic People's Republic of Korea	SD	Sudan
CF	Central African Republic	KR	Republic of Korea	SE	Sweden
CG	Congo	KZ	Kazakhstan	SG	Singapore
CH	Switzerland	LI	Liechtenstein	SI	Slovenia
CI	Côte d'Ivoire	LK	Sri Lanka	SK	Slovakia
CM	Cameroon	LR	Liberia	SN	Senegal
CN	China	LT	Lithuania	SZ	Swaziland
CS	Czechoslovakia	LU	Luxembourg	TD	Chad
CZ	Czech Republic	LV	Latvia	TG	Togo
DE	Germany	MC	Monaco	TJ	Tajikistan
DK	Denmark	MD	Republic of Moldova	TT	Trinidad and Tobago
EE	Estonia	MG	Madagascar	UA	Ukraine
ES	Spain	ML	Mali	UG	Uganda
FI	Finland	MN	Mongolia	US	United States of America
FR	France	MR	Mauritania	UZ	Uzbekistan
GA	Gabon			VN	Viet Nam

-1-

DESCRIPTION**STABLE PRESERVATION METHOD OF POWDERED SOFT DRINK PREPARATION
AND POWDERED SOFT DRINK PREPARATION****TECHNICAL FIELD**

5 The present invention relates to a stable
preservation method of a powdered soft drink preparation, and
a powdered soft drink preparation.

BACKGROUND ART

10 A powdered soft drink preparation normally contain a
carbohydrate and a sour agent as a main component, and is
prepared by additionally adding a subraw material such as
fruit juice. It is necessary that the powdered soft drink
preparation maintains the dried state as much as possible
because of it's high moisture absorption. That is, the soft
15 drink preparation causes deterioration such as coagulation,
browning, discoloration in the presence of water, which
results in deterioration of the quality.

20 In addition, this preparation always contain about 1
% of water derived from the raw material even if water is
severely controlled. When the component causing the reaction
between water and the above main component is present in the
subraw material, for example, when a basic nutritive component
is present, the neutralization reaction proceeds with time.
When amino acid is added, the browning due to Mailard reaction
25 arises. Therefore, there is a drawback in both cases that it

-2-

becomes very difficult to maintain the preparation stably.

Furthermore, even if the preparation is reserved in an enclosed container, it is difficult to avoid deterioration of quality and change of taste with time, which are caused by the above-described water of about 1 %, water adhered on raw materials, water to be formed by the neutralization reaction between the added components. Therefore, it is essential for the preparation to use a desiccant at present. Thus, it is requested to develop a novel method for preventing deterioration due to water for a long period of time to maintain the quality.

DISCLOSURE OF THE INVENTION

A main object of the present invention is to solve a drawback such as deterioration caused in the presence of water, which is admitted in a conventional powdered soft drink preparation, thereby providing a powdered soft drink having an excellent preservation stability for a long period of time.

The present inventors have intensively studied so as to accomplish the above object. As a result, it has been found that a powdered soft drink preparation having an excellent preservation stability can be obtained when a predetermined amount of a calcium oxide and a particulate silicon dioxide are added in a conventional powdered soft drink preparation.

That is, according to the present invention, there

-3-

is provided a stable preservation method of a powdered soft drink preparation, which comprises adding 0.2 to 1.0 % by weight of a calcium oxide and 0.02 to 2.0 % by weight of a particulate silicon dioxide in the powdered soft drink

5 preparation containing a carbohydrate and a sour agent as a main component. There is also provided a powdered soft drink preparation thus obtained and having an improved reservation stability.

According to the method of the present invention, it
10 is considered that the calcium oxide to be added reacts chemically with water derived from the raw material in the preparation to remove water. Thereby, a water content in the preparation is reduced to stabilize the preparation. As a result, the preparation exhibits a high preservation
15 stabilization effect for a sufficiently long period of time without using no desiccant. On the other hand, it is considered that the particulate silicon dioxide serves as a so-called coating agent which coats the surfaces of the raw material powders to reduce the opportunity of direct
20 contacting between powders having a high reactivity, thereby improving physical properties and stability of the preparation.

According to the method of the present invention, there can be obtained a powdered soft drink preparation having
25 a high preservation stability which has never been

accomplished, heretofore. That is, regarding the preparation of the present invention, the commercial value such as flavor is not deteriorated, and drawbacks such as coagulation and browning are prevented. Furthermore, the above-mentioned components to be added have safety guaranteed in the field of the food industry. At the same time, they are tasteless and odorless and, therefore, there is not any fear of exerting a harmful influence on the taste of the soft drink preparation itself.

10 BEST MODE FOR CARRYING OUT THE INVENTION

The calcium oxide to be added in the powdered soft drink preparation is contained in an amount of not less than 0.2 % by weight, thereby exhibiting the desired stabilization effect of the preparation. Even if the large amount of the calcium oxide is added, the stabilization effect is not improved in proportion to the amount, and the taste, flavor, solubility, etc. of the preparation are likely to be deteriorated. Therefore, it is preferred to use the calcium oxide in an amount of not more than 1.0 % by weight. Particularly preferred amount is selected within a range of 0.2 to 0.5 % by weight.

Regarding the silicon dioxide, the desired effect can be exhibited by adding in an amount within a range of 0.02 to 2.0 % by weight, and it is preferred to use it in an amount within a range of 0.03 to 1.0 % by weight.

-5-

The calcium oxide is not specifically limited, but those synthesized chemically and those obtained by calcining shells or egg shells derived from natural products can be advantageously used in the present invention. Among them, the
5 chemically synthesized calcium oxide and shell-calcined calcium derived from natural products are particularly preferred from the viewpoint of organoleptic properties such as taste of the resulting preparation.

The particulate silicon dioxide is not also
10 specifically limited and various ones which are known to public can be used. Among them, those having a three-dimensional amorphous structure like silica gel is normally preferred. Further, as the particulate silicon dioxide, those having an average particle size of not more than 15 μm , which
15 are defined as a food additive at present, are used.

The powdered soft drink preparation to be stabilized by adding the above two specific components according to the present invention may be the same composition as that of a most popular one containing, as a main component, a
20 carbohydrate and an organic acid as a sour agent. The amount of the main component to be added can be decided within a normal range according to the shape, use and purpose of using of the preparation. It is preferred that the amount of the carbohydrate in order to obtain good organoleptic properties
25 is selected within a range of about 65 to 95 % by weight in

-6-

the powdered preparation. After the carbohydrate was dissolved in water, the amount is selected within a range of about 4 to 12 % by weight.

Examples of the above carbohydrate are one or more of normal carbohydrates, which can be selected from monosaccharides such as glucose and fructose; disaccharides such as sucrose and maltose; polysaccharides such as oligosaccharide. The effect of the present invention is enhanced, when using monosaccharides (e.g. fructose) which have high moisture absorption and high reactivity.

The sour agent to be used as another main component may be various organic acids which are normally used. Typical examples include citric acid, tartaric acid, fumaric acid, malic acid, lactic acid and ascorbic acid. They may be used alone or in combination.

Optional components such as various vitamins, amino acids and minerals can be appropriately added in the powdered soft drink preparation of the present invention according to the purpose of using. In addition to them, there can be appropriately added binders, excipients, disintegrators, lubricants, thickeners, surfactants, osmotic pressure adjustors, electrolytes, sweeteners, flavors, pigments, pH adjustors and the like.

The powdered soft drink preparation of the present invention can be prepared according to the same manner as in

-7-

the normal powdered soft drink preparation except for adding a predetermined amount of the above two components. The time of the addition of the above essential two components is not specifically limited and it may be any time in the desired preparation step of the preparation.

Thus, the desired powdered soft drink preparation whose preservation stability has been improved can be obtained.

FIELD OF INDUSTRIAL APPLICABILITY

According to the method of the present invention, a preservation stability of a powdered soft drink preparation is remarkably improved. As a result, regarding the preparation of the present invention, the commercial value such as flavor is not deteriorated, and drawbacks such as the coagulation and browning are prevented. Components to be added as preservative stabilizers have safety guaranteed in the field of the food industry. At the same time, they are tasteless and odorless and, therefore, there is not any fear of exerting a harmful influence on the taste of the soft drink preparation itself.

EXAMPLES

In order to explain the present invention in detail, Preparation Examples of the preparation of the present invention will be described below. Then, Stability Test carried out with respect to the resulting preparation will be

described. In the respective Examples, "percentages" are by weight unless otherwise stated.

Examples 1 to 9

5 The respective components described in Table 1 were weighed and mixed to prepare powdered soft drink preparations. These are dissolved in 1 liter of water to be served as a drink.

10

15

20

25

Table 1

Component (%)	Examples No.								
	1	2	3	4	5	6	7	8	9
Glucose	-	-	-	80	80	80	-	-	-
Fructose	80	80	80	-	-	-	-	-	-
Granulated sugar	-	-	-	-	-	-	80	80	80
Sour agent	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.
L-ascorbic acid	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0	3.0
L-arginine	3.3	3.3	3.3	3.3	3.3	3.3	3.3	3.3	3.3
Shell-calcined calcium	0.25	0.5	1.0	0.25	0.5	1.0	0.25	0.5	1.0
Particulate silicon dioxide	0.05	0.1	0.5	0.05	0.1	0.5	0.05	0.1	0.5
Powdered juice	2.0	1.5	1.0	2.0	1.5	1.0	2.0	1.5	1.0
Various vitamins	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.
Flavor	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.	s.a.
Total weight (g)	62.5	62.5	62.5	62.5	62.5	62.5	62.5	62.5	62.5
s.a.: suitable amount									

Preservation Stability Test 1

The preparation of the present invention obtained according to the following Preparation Example I and the following control preparations were subjected to a comparison test.

Preparation Example A (formulation)

Fructose	8.3 %
Sour agent	suitable amount
L-arginine	3.3 %
10 L-ascorbic acid	2.5 %
Powdered fruit juice	2.0 %
Sodium chloride	suitable amount
Shell-calcined calcium	0.25 %
Particulate silicon dioxide	0.05 %
15 Flavor and pigment	suitable amount
Total	100 % (total amount: 62.5 g)

(1) Control preparation (containing no shell-calcined calcium): preparation obtained according to the same manner as that described in Preparation Example A except for adding no shell-calcined calcium

(2) Control product 1: preparation obtained according to the same manner as that described in Preparation Example A except for adding no shell-calcined calcium and particulate silicon dioxide

25 (3) Control product 2: preparation obtained according to the

-11-

same manner as that described in Preparation Example A except for adding no particulate silicon dioxide

The above respective preparations were prepared by mixing with hands and packed separately in a bag made of an aluminum laminated film, respectively. The bag was preserved in a constant temperature room and the preservation stability thereof was tested.

The followings were selected as the test item.

1. Coagulation of preparation
2. Degree of discoloration of preparation
3. Change in organoleptic property (taste) in dissolution of preparation

The results were as follows.

1. Coagulation of preparation

The preparation of Preparation Example A (product of the present invention) and control preparation (containing no shell-calcined calcium, controls 1 and 2) were allowed to stand in a constant temperature room at 50 °C to visually observe the state of the respective samples after 2, 3 and 4 weeks, respectively.

As a result, the product of the present invention caused no coagulation even after standing at 50 °C for 4 weeks and was superior in powder fluidity. To the contrary, in all control preparations, the coagulation state became serious by adding no shell-calcined calcium and/or particulate silicon

-12-

dioxide. After standing for 3 weeks, the preparations were coagulated tightly and showed a sheet-like state (regarding the control product 2, the mass was sometimes broken because the degree of the coagulation is slightly low) and it could not be easily returned to the powder state. Furthermore, the solubility of the respective control preparations in water became inferior, in comparison with the preparation of the present invention after standing under the same condition.

2. Degree of discoloration of preparation

The respective preparation samples whose coagulation state was judged according to the above item 1 were visually observed and the change in color with time of the respective samples was examined using a color difference meter. Further, the value after a predetermined time has been passed was judged by NBS unit (ΔE : color difference) and Lab, using the value before the beginning of the test as a standard. The explanation of the value of the NBS unit and indication of visual sense ("Matter with respect to Color", published by Nihon Denshoku Kogyo) as well as evaluation of Lab are as follows.

(NBS unit)

0 to 0.5

0.5 to 1.5

1.5 to 3.0

(Visual sense)

faintly

slightly

in the degree capable of
perceiving

-13-

3.0 to 6.0	in the degree of being
	remarkable
6.0 to 12.0	greatly
12.0 or more	drastically

5 Lab

L: The larger the value of L, the higher the brightness is.

a: It shows a degree of red at the (+) side, and a degree of green at the (-) side.

10 b: It shows a degree of yellow at the (+) side, and a degree of blue at the (-) side.

The test results of the preparation of the Preparation Example A (product of the present invention) and control product 1 are shown in Tables 2 and 3.

15

20

25

Table 2

Condition of change with time (NBS)	Preparation A (product of (the present invention)	Control product 1
50 °C, 2 weeks	2.38	15.00
50 °C, 3 weeks	4.70	16.20
50 °C, 4 weeks	5.01	23.19

- 15 -

Table 3

Condition of change with time (Lab)	Preparation A (product of (the present invention))		Control product 1	
	L	a b	L	a b
at the time of beginning	89.62	0.93 6.74	89.68	1.21 5.84
50 °C, 2 weeks	89.23	0.03 8.91	80.33	2.54 18.41
50 °C, 3 weeks	88.29	1.05 11.25	78.65	2.84 18.51
50 °C, 4 weeks	88.43	0.28 11.56	71.03	4.89 20.03

-16-

As is apparent from the above tables, the discoloration was hardly observed in the product of the present invention, in comparison with the control product 1. That is, regarding the product of the present invention, the
5 brightness hardly changed and a stable color tone was shown even after preservation at 50 °C for 4 weeks. To the contrary, regarding the control product 1, the brightness was decreased with time during preserving at 50 °C and the degree of red and yellow was increased. This means that the fact.
10 that browning proceeds with time is showed clearly with the numerical value.

Actually, regarding the product of the present invention, little discoloration was admitted by visual observation even after preservation at 50 °C for 4 weeks. To
15 the contrary, regarding the control product 1, a remarkable discoloration has already been admitted after 2 weeks.

Further, regarding the control preparation containing no shell-calcined calcium and the control product 2, a remarkable discoloration which is almost the same as that
20 of the above control product 1 was admitted. Among them, the control product 2 tends to cause a weak degree of browning.

3. Change in organoleptic property (taste) in dissolution of preparation

A standard sample which was preserved in a dark
25 place at 4°C after preparation, and samples which were

-17-

obtained by preserving the standard samples at 37°C or 50°C for a predetermined time were used. The change in taste of each sample due to preservation was evaluated by asking three panelists to try each drink which was prepared by dissolving
5 62.5 g of each sample in 1 liter of cold water. Evaluation was carried out on the basis of the following five-level marks in comparison of the standard sample by the organoleptic test.

5 Marks: Not changed.

4 Marks: The flavor became faint, slightly, but the total
10 unity of the taste was not changed.

3 Marks: The flavor became faint and the pleasant taste was disappeared.

2 Marks: The change in flavor and taste were clearly admitted and the nasty smell was also admitted.

15 1 Mark: The flavor becomes inferior, drastically, and the nasty taste and smell (e.g. bitter taste, burnt smell, etc.) were admitted.

The results of the product of the present invention and control 1 are shown in the following Table 4.

20

25

Table 4

Condition of change with time (Lab)	Preparation A (product of (the present invention)	Control product 1
50 °C, 2 weeks	4 marks	2 marks
50 °C, 3 weeks	3 marks	1 mark
50 °C, 4 weeks	3 marks	1 mark
37 °C, 1 month	5 marks	3 marks
37 °C, 2 months	4 marks	2 marks

-19-

As is apparent from Table 4, regarding the product of the present invention, the change in taste did not become a problem, substantially, even after preservation at 50 °C for 2 weeks (37 °C, 2 months). To the contrary, regarding the control 1, the taste has already changed after preservation at 37 °C for one month. After preservation at 50 °C for 2 weeks, an clear change in taste was admitted and an initial pleasant flavor was disappeared organoleptically. Therefore, a main flavor was replaced by a burnt smell by which the browning is accompanied, which results in flavor which is far different from that of a normal soft drink.

Further, regarding the control preparation containing no shell-calcined calcium and control 2, a clear change in taste could be observed, similar to the above control product 1.

CLAIMS

1. A stable preservation method of a powdered soft drink preparation, which comprises adding 0.2 to 1.0 % by weight of a calcium oxide and 0.02 to 2.0 % by weight of a particulate silicon dioxide in the powdered soft drink preparation containing a carbohydrate and a sour agent as a main component, each amount being based on the total amount of the preparation.
2. A stable preservation method according to claim 1, wherein the calcium oxide is selected from a chemically synthesized calcium oxide and a shell-calcined calcium derived from a natural product.
3. A stable preservation method according to claim 1, wherein the particulate silicon dioxide has a three-dimensional amorphous structure.
4. A stable preservation method according to claim 1, wherein an average particle size of the particulate silicon dioxide is not more than 15 μm .
5. A stabilized powdered soft drink preparation, comprising a carbohydrate and a sour agent as a main component, wherein 0.2 to 1.0 % by weight of a calcium oxide and 0.02 to 2.0 % by weight of a particulate silicon dioxide are added as a preservation stabilizer.
6. A powdered soft drink preparation according to claim 5, wherein the carbohydrate is added in a proportion of

-21-

65 to 95 % by weight, based on the total amount of the powder.

5

10

15

20

25

INTERNATIONAL SEARCH REPORT

International Application No

PCT/96/00109

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 A23L2/44 A23L2/39

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 A23L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	GB,A,2 183 808 (HAYASHIBARA) 10 June 1987 see page 1, line 11-17 ---	1
A	US,A,4 278 695 (V.S.VELASCO) 14 July 1981 see column 2, line 14-24; claim; example ---	1,4
A	DATABASE WPI Section Ch, Week 8346 Derwent Publications Ltd., London, GB; Class D13, AN 83-817418 XP002002267 & JP,A,58 170 464 (NISSAN) see abstract --- -/--	1

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- *&* document member of the same patent family

Date of the actual completion of the international search

7 May 1996

Date of mailing of the international search report

15.05.96

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
 NL - 2280 HV Rijswijk
 Tel. (+ 31-70) 340-2040, Tx. 31 651 epo nl,
 Fax: (+ 31-70) 340-3016

Authorized officer

Van Moer, A

INTERNATIONAL SEARCH REPORT

International Application No

JP 96/00109

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>DATABASE WPI Section Ch, Week 9031 Derwent Publications Ltd., London, GB; Class A92, AN 90-236040 XP002002268 & JP,A,02 164 536 (ASAHI) see abstract -----</p>	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/JP96/00109

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB-A-2183808	10-06-87	JP-B- 7010341	08-02-95
		JP-A- 62125854	08-06-87
		JP-C- 1745069	25-03-93
		JP-B- 4029415	18-05-92
		JP-A- 62136240	19-06-87
		AU-B- 598468	28-06-90
		BE-A- 905157	23-01-87
		CA-A- 1296176	25-02-92
		DE-A- 3625611	04-06-87
		FR-A- 2590500	29-05-87
		US-A- 4996196	26-02-91
		US-A- 4870059	26-09-89
		KR-B- 9400165	08-01-94

US-A-4278695	14-07-81	CA-A- 1132834	05-10-82
